

OCT - 9 2001

**510(k) Summary**  
(As required by 21 CFR 807.92)**A. Submitter Information**

Submitter's Name: St. Jude Medical, Daig Division  
Address: 14901 DeVeau Place  
Minnetonka, Minnesota 55345-2126 U.S.A.  
Telephone Number: (952) 352-9769  
Contact Person: Jim M. Taufen  
Date Submission Prepared: 07-Sep-2001

**B. Device Information**

Trade Name: Seal-Away CS™ Introducer Kit  
Common or Usual Name: Percutaneous Catheter Introducer  
Classification Name: Percutaneous Catheter Introducer (per 21 CFR Part 870.1340)  
Device Classification: Class II (per 21 CFR Part 870.1340)  
Panel - Cardiovascular  
Predicate Device: Peel away Sheath Introducer and Obturator  
St. Jude Medical, Daig Division  
Device Description: The Seal-Away CS™ introducer kit includes a Peel-Away Introducer, Obturator, Seal-Away™ detachable hemostasis valve, Guidewire, Syringe and Needle. The introducer kits are provided sterile, and are intended for single-use only.  
Intended Use: The Seal-Away CS Introducer kits are designed for the introduction of catheters into a vessel.

**C. Comparison of Required Technological Characteristics**

All technological characteristics of the Seal-Away™ CS introducer kit are substantially equivalent to the predicate device including product design, packaging, sterilization, and labeling.

**D. Support of the Substantial Equivalence**

St. Jude Medical, Daig Division considers the Seal-Away CS introducer kit to be substantially equivalent to the predicate device.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

OCT - 9 2001

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Jim M. Taufen  
St. Jude Medical  
Daig Division  
14901 DeVeau Place  
Minnetonka, MN 55345

Re: K013029  
Seal-Away CS™ Introducer Kit  
Regulation Number: 870.1340  
Regulation Name: Catheter Introducer  
Regulatory Class: Class II  
Product Code: DYB  
Dated: September 7, 2001  
Received: September 10, 2001

Dear Mr. Taufen:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with

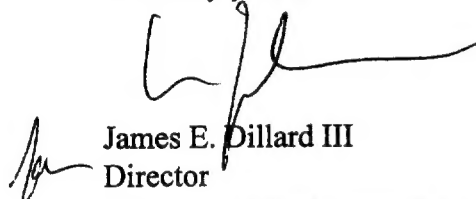
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all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4586. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



James E. Dillard III  
Director  
Division of Cardiovascular and  
Respiratory Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure

510(k) Number (if known): K013029

Device Name: Seal-Away CS™ Introducer Kit

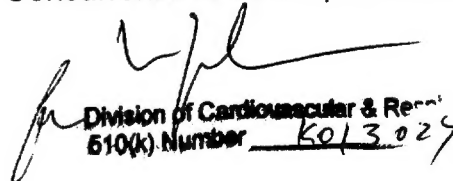
Indications for Use:

The Seal-Away CS™ Introducer Kit is a percutaneous catheter introducer kit with peel-away sheath that is indicated for the introduction of catheters into a vessel.

The Seal-Away™ Hemostasis Adapter is indicated for the introduction of pacing leads, angiographic catheters, closed end catheters, balloon catheters and electrodes into a vessel where minimizing blood loss is essential.

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

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Concurrence of CDRH, Office of Device Evaluation (ODE)

  
Division of Cardiovascular & Renal  
510(k) Number K013029

Prescription Use ☒  
(Per 21 CFR 801.109)

OR

Over-The-Counter Use \_\_\_\_\_  
(Optional Format 1-2-96)